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(54) Title: 2-AMINOQUINOLINECARBOXAMIDES: NEUROKININ RECEPTOR LIGANDS			
(57) Abstract			
<p>Disclosed are compounds of formula (I) or pharmaceutically acceptable salts or pharmaceutically acceptable solvates thereof, wherein R₁, R₂, R₃, X, Q₁ and Q₂ are defined herein, which compounds are ligands for neurokinin receptors, in particular NK-3 receptors, and are therefore useful in the treatment of a wide range of diseases or disorders including, but not limited to depression, anxiety, psychosis, obesity, pain, Parkinson's disease, Alzheimer's disease, neurodegenerative diseases, movement disorders, respiratory diseases, inflammatory diseases, neuropathy, immune disorders, migraine, biliary disfunction, and dermatitis.</p>			
<div style="text-align: right;">(I)</div>			

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2-AMINOQUINOLINECARBOXAMIDES:
NEUROKININ RECEPTOR LIGANDS

5 This application claims priority from provisional application no. 60/131,025 which is incorporated herein by reference in its entirety.

BACKGROUND OF THE INVENTION

10

Field of the Invention

 This invention relates to 2-aminoquinolinecarboxamides, pharmaceutical compositions comprising them, and the use of such compounds in the treatment of certain central nervous system and
15 peripheral diseases. The compounds of this invention are also useful as probes for the localization of cell surface receptors.

Description of the related art

 The tachykinins represent a family of structurally related
20 peptides originally isolated based upon their smooth muscle contractile and sialogogic activity. These mammalian peptides include substance P (SP), neurokinin A (NKA) and neurokinin β (NKB). Tachykinins are synthesized in the central nervous system (CNS), as well as in peripheral tissues, where they exert a
25 variety of biological activities. Substance P can be produced from three different mRNAs (α -, β - and γ -preprotachykinin mRNAs) that arise from a single gene as a result of alternative RNA splicing, whereas NKA can be generated from either the β - or the γ -preprotachykinin mRNA through posttranslationally processed
30 precursor polypeptides. These precursors can also be differentially processed so that amino terminally extended forms of NKA (known as neuropeptide K and neuropeptide γ) are produced. NKB is produced from a separate mRNA arising from a second gene known as preprotachykinin B.

35 Three receptors for the tachykinin peptides have been molecularly characterized and are referred to as neurokinin-1

(NK-1), neurokinin-2 (NK-2) and neurokinin-3 (NK-3) receptors. The NK-1 receptor has a natural agonist potency profile of SP>NKA>NKB. The NK-2 receptor agonist potency profile is NKA>NKB>SP, and the NK-3 receptor agonist potency profile is NKB>NKA>SP. These receptors mediate the variety of tachykinin-stimulated biological effects that generally include 1) modulation of smooth muscle contractile activity, 2) transmission of (generally) excitatory neuronal signals in the CNS and periphery (e.g. pain signals), 3) modulation of immune and inflammatory responses, 4) induction of hypotensive effects via dilation of the peripheral vasculature, and 5) stimulation of endocrine and exocrine gland secretions. These receptors transduce intracellular signals via the activation of pertussis toxin-insensitive ($G_{\alpha q/11}$) G proteins, resulting in the generation of the intracellular second messengers inositol 1,4,5-trisphosphorylate and diacylglycerol. NK-1 receptors are expressed in a wide variety of peripheral tissues and in the CNS. NK-2 receptors are expressed primarily in the periphery, while NK-3 receptors are primarily (but not exclusively) expressed in the CNS. Recent work confirms the presence of NK-3 receptor binding sites in the human brain.

Studies measuring the localization of NKB and NK-3 receptor mRNAs and proteins, along with studies performed using peptide agonists and non-peptide NK-3 receptor antagonists, provide a rationale for using NK-3 receptor antagonists in treating a variety of disorders in both the CNS and the periphery. In the CNS, activation of NK-3 receptors has been shown to modulate dopamine and serotonin release, indicating therapeutic utility in the treatment of a variety of disorders including anxiety, depression, schizophrenia and obesity. Further, studies in primate brain detect the presence of NK-3 mRNA in a variety of regions relevant to these disorders. With regard to obesity, it has also been shown that NK-3 receptors are located on MCH-containing neurons in the rat lateral hypothalamus and zona

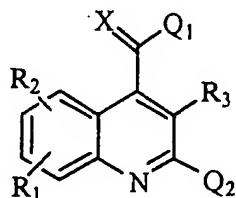
incerta. In the periphery, administration of NKB into the
airways is known to induce mucus secretion and
bronchoconstriction, indicating therapeutic utility for NK-3
receptor antagonists in the treatment of patients suffering from
5 airway diseases such as asthma and chronic obstructive pulmonary
disease (COPD). Localization of NK-3 receptors in the
gastrointestinal (GI) tract and the bladder indicates
therapeutic utility for NK-3 receptor antagonists in the
treatment of GI and bladder disorders including inflammatory
10 bowel disease and urinary incontinence.

Both peptide and nonpeptide antagonists have been developed
for each of the tachykinin receptors. The first generation of
peptide antagonists for the tachykinin receptors had problems
with low potency, partial agonism, poor metabolic stability and
15 toxicity, whereas the current generation of non-peptide
antagonists display more drug-like properties. Unfortunately,
previous non-peptide NK-3 receptor antagonists suffer from a
number of problems such as species selectivity (which limits the
potential to evaluate these compounds in many appropriate
20 disease models). New non-peptide NK-3 receptor antagonists are
therefore being sought, both as therapeutic agents and as tools
to further investigate the anatomical and ultrastructural
distribution of NK-3 receptors, as well as the physiologic and
pathophysiologic consequences of NK-3 receptor activation.

25

SUMMARY OF THE INVENTION

This invention relates to 2-aminoquinolinecarboxamides represented by structure Formula I:



Formula I

or pharmaceutically acceptable salts or pharmaceutically acceptable solvates thereof, wherein in R₁, R₂, R₃, X, Q₁ and Q₂ are hereinafter defined.

The invention also relates to pharmaceutical compositions comprising such compounds and the use of such compounds and compositions in the treatment of certain central nervous system and peripheral diseases or disorders.

The compounds of this invention are ligands for neurokinin receptors, in particular NK-3 receptors, and are useful in the treatment of a wide range of diseases or disorders including, but not limited to depression, anxiety, psychosis, obesity, pain, Parkinson's disease, Alzheimer's disease, neurodegenerative diseases, movement disorders, respiratory diseases, inflammatory diseases, neuropathy, immune disorders, migraine, biliary disfunction, and dermatitis.

The invention provides novel 2-aminoquinolinecarboxamides that bind selectively to neurokinin receptors, in particular NK-3 receptors. These compounds are therefore of use in the treatment of a broad array of diseases which are characterized by modulation of the neurokinin receptors, in particular NK-3 receptors.

In a separate aspect, the invention provides methods of using compounds of this invention as positive controls in assays

for receptor activity and using appropriately labeled compounds of the invention as probes for the localization of receptors, particularly neurokinin-3 receptors, in tissue sections.

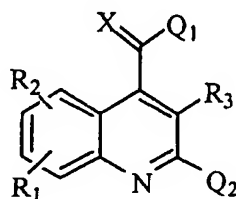
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DETAILED DESCRIPTION OF THE INVENTION

This invention relates to 2-aminoquinolinecarboxamides, pharmaceutical compositions comprising them, and the use of such compounds in the treatment of certain central nervous system and peripheral diseases or disorders.

10

Accordingly, a broad embodiment of the invention is directed to compounds of Formula I:



I

or pharmaceutically acceptable salts or pharmaceutically acceptable solvates thereof, wherein:

15

R₁ is:

hydrogen, halogen, hydroxy, C₁₋₆ alkyl, -O(C₁₋₆ alkyl), -NO₂,
 -CN, -SO₂NH₂, -SO₂NH(C₁₋₆ alkyl), -SO₂N(C₁₋₆ alkyl)(C₁₋₆ alkyl),
 20 amino, -NH(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)(C₁₋₆ alkyl), -N(C₁₋₆
 alkyl)CO(C₁₋₆ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)CO₂(C₁₋₆
 alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)SO₂(C₁₋₆ alkyl),
 -SO₂NHCO(C₁₋₆ alkyl), -CONHSO₂(C₁₋₆ alkyl), -CON(C₁₋₆ alkyl)
 (C₁₋₆ alkyl), -CO₂(C₁₋₆ alkyl), -S(C₁₋₆ alkyl), -SO(C₁₋₆ alkyl),
 25 or -SO₂(C₁₋₆ alkyl),

25

where each C₁₋₆ alkyl is a straight, branched or cyclic alkyl group optionally containing one or two double or triple bonds and optionally substituted with one or more of hydroxy, oxo, halogen, amino, or C₁₋₃ alkoxy;

30

R₂ and R₃ are independently selected from the groups consisting of:

hydrogen, halogen, hydroxy, C₁₋₈ alkyl, -O(C₁₋₈ alkyl), -NO₂,
 -CN, -SO₂NH₂, -SO₂NH(C₁₋₈ alkyl), -SO₂N(C₁₋₈ alkyl)(C₁₋₈ alkyl),
 5 amino, -NH(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)(C₁₋₈ alkyl), -N(C₁₋₈
 alkyl)CO(C₁₋₈ alkyl), -NHCO₂(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)CO₂(C₁₋₈
 alkyl), -NHSO₂(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)SO₂(C₁₋₈ alkyl), -
 SO₂NHCO(C₁₋₈ alkyl), -CONHSO₂(C₁₋₈ alkyl), -CON(C₁₋₈ alkyl)(C₁₋₈
 alkyl), -CO₂(C₁₋₈ alkyl), -S(C₁₋₈ alkyl), -SO(C₁₋₈ alkyl), -
 10 SO₂(C₁₋₈ alkyl), and Ar,

wherein Ar is phenyl, naphthyl, thienyl, benzothienyl,
 pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl,
 benzoimidazolyl, furanyl, benzofuranyl, thiazolyl,
 benzothiazolyl, isothiazolyl, benzisothiazolyl, triazolyl,
 15 tetrazolyl, pyrazolyl, or benzopyrazolyl, each of which
 is optionally substituted with one or more of:

halogen, hydroxy, C₁₋₆ alkyl, -O(C₁₋₆ alkyl), -NO₂, -CN, -
 SO₂NH₂, -SO₂NH(C₁₋₆ alkyl), -SO₂N(C₁₋₆ alkyl)(C₁₋₆ alkyl),
 amino, -NH(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)(C₁₋₆ alkyl), -N(C₁₋₆
 20 alkyl)CO(C₁₋₆ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₆
 alkyl)CO₂(C₁₋₆ alkyl), -NHSO₂(C₁₋₆ alkyl), -N(C₁₋₆
 alkyl)SO₂(C₁₋₆ alkyl), -SO₂NHCO(C₁₋₆ alkyl), -CONHSO₂(C₁₋₆
 alkyl), -CON(C₁₋₆ alkyl)(C₁₋₆ alkyl), -CO₂(C₁₋₆ alkyl), -
 S(C₁₋₆ alkyl), -SO(C₁₋₆ alkyl), or -SO₂(C₁₋₆ alkyl),

25 where

each C₁₋₆ alkyl is as defined above for R₁; and
 each C₁₋₈ alkyl is a straight, branched or cyclic alkyl group
 optionally containing one or two double or triple bonds and
 optionally substituted with one or more of:

- 30 (i) hydroxy,
 (ii) oxo,
 (iii) halogen,
 (iv) Ar, wherein Ar is as defined above,

(v) $-NR_4R_5$, wherein R_4 and R_5 are independently selected from:

(A) hydrogen;

(B) C_{1-6} alkyl, wherein C_{1-6} alkyl is as defined above for R_1 ;

(C) Ar, where in Ar is as defined above;

(D) C_{1-6} alkyl-Ar, wherein Ar is as defined above and Ar is attached to any position of the C_{1-6} alkyl group at any position of Ar;

or R_4 and R_5 together form a 4- to 8-membered monocyclic or bicyclic nitrogen-containing ring which may contain:

(a) one or two double bonds;

(b) one or two oxo;

(c) one or two of O, S or $N-R_6$ wherein R_6 is hydrogen, C_{1-6} alkyl, C_{1-6} alkyl-Ar, wherein C_{1-6} alkyl and Ar are defined as above and Ar is attached to to any position of the C_{1-6} alkyl group at any position of Ar; or

(d) one or two of R_1 where R_1 is as defined above;

(vi) $-OR_4$, wherein R_4 is as defined above;

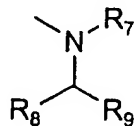
(vii) $-CONR_4R_5$, wherein R_4 and R_5 are as defined above;

(viii) $-SO_2NR_4R_5$, wherein R_4 and R_5 are as defined above;

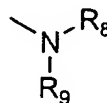
(x) $-NR_4COR_5$, wherein R_4 and R_5 are as defined above;

X is O, S or N-CN;

Q_1 and Q_2 are independently selected from formulas II and III:



II



III

wherein

R_7 is hydrogen or C_{1-8} alkyl;

C₁₋₈ alkyl is as defined above;

R₈ and R₉ are independently selected from:

(i) hydrogen;

(ii) Ar', wherein Ar' is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, benzoimidazolyl, furanyl, benzofuranyl, thiazolyl, benzothiazolyl, isothiazolyl, benzisothiazolyl, triazolyl, tetrazolyl, pyrazolyl, or benzopyrazolyl, each of which is unsubstituted or substituted with one or more of:

halogen, hydroxy, C₁₋₈ alkyl, -O(C₁₋₈ alkyl), -NO₂, -CN, -SO₂NH₂, -SO₂NH(C₁₋₈ alkyl), -SO₂N(C₁₋₈ alkyl)(C₁₋₈ alkyl), amino, -NH(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)CO(C₁₋₈ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₈ alkyl)CO₂(C₁₋₈ alkyl), -NHSO₂(C₁₋₈ alkyl), -N(C₁₋₆ alkyl)SO₂(C₁₋₈ alkyl), -SO₂NHCO(C₁₋₈ alkyl), -CONHSO₂(C₁₋₈ alkyl), -CON(C₁₋₈ alkyl)(C₁₋₈ alkyl), -CO₂(C₁₋₈ alkyl), -S(C₁₋₈ alkyl), -SO(C₁₋₈ alkyl), or -SO₂(C₁₋₈ alkyl), wherein C₁₋₈ alkyl and C₁₋₆ alkyl are as defined above;

(iii) C₁₋₈ alkyl, wherein said C₁₋₈ alkyl is as defined above,

(iv) -COR₄, wherein R₄ is as defined above,

(v) -CONR₄R₅, wherein R₄ and R₅ are as defined above,

(vi) -SO₂NR₄R₅, wherein R₄ and R₅ are as defined above,

or R₈ and R₉ together form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

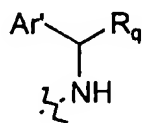
(a) one or two double bonds;

(b) one or two oxo;

(c) one or two O, S or N-R₁₀ wherein R₁₀ is hydrogen, C₁₋₈ alkyl, C₁₋₈ alkyl-Ar', wherein C₁₋₈ alkyl and Ar' are defined as above and Ar' may be attached to C₁₋₈ alkyl at any position; or

(d) one or two R_1 groups.

Preferred compounds of the invention include those of Formula I where Q_1 is a group of the formula I-g:



I-g

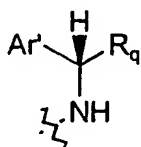
where

Ar' is phenyl optionally substituted with one or more of halogen, hydroxy, C_{1-8} alkyl, $-O(C_{1-8}$ alkyl), $-\text{NO}_2$, $-\text{CN}$, amino, $-\text{NH}(C_{1-8}$ alkyl), $-\text{N}(C_{1-8}$ alkyl)(C_{1-8} alkyl), $-\text{N}(C_{1-8}$ alkyl) $\text{CO}(C_{1-8}$ alkyl), $-\text{NHCO}_2(C_{1-6}$ alkyl), $-\text{N}(C_{1-8}$ alkyl) $\text{CO}_2(C_{1-8}$ alkyl), $-\text{CON}(C_{1-8}$ alkyl)(C_{1-8} alkyl), $-\text{CO}_2(C_{1-8}$ alkyl), wherein C_{1-8} alkyl and C_{1-6} alkyl are as defined above; and

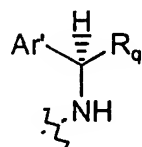
R_q is straight or branched chain alkyl having from 1-6 carbon atoms.

More preferred compounds of Formula I where Q_1 is a group of formula I-g include those where X is oxygen. Particularly preferred are such compounds where R_1 and R_2 are both hydrogen.

Formula I-g encompasses the following groups:



I-h

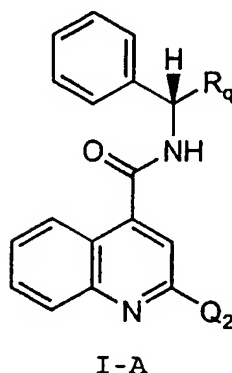


I-k

where Ar' and R_q carry the same definitions as given above for formula I-g.

The invention encompasses compounds having Q_1 groups represented by either of formulas I-h or I-k.

Other preferred compounds of Formula I include those of Formula I-A:



5 where R_q is straight or branched chain alkyl having from 1-6 carbon atoms; and

Q_2 is as defined above for Formula I.

Particularly preferred compounds of Formula I-A are those where R_q is ethyl; Q_2 represents formula I or formula II where R_1 is defined as above and R_8 and R_9 are independently selected from:

hydrogen;

C_{1-8} alkyl, wherein said C_{1-8} alkyl is as defined above; or

15 R_8 and R_9 together form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

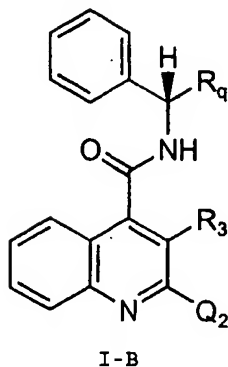
(a) one or two double bonds;

(b) one or two oxo;

(c) one or two O, S or N- R_{10} wherein R_{10} is hydrogen, C_{1-8} alkyl, C_{1-8} alkyl-Ar', wherein C_{1-8} alkyl and Ar' are defined as above and Ar' may be attached to C_{1-8} alkyl at any position; or

(d) one or two R_1 groups.

25 Still other preferred compounds of Formula I include those of Formula I-B:



Wherein:

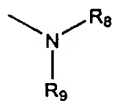
R_q is straight or branched chain alkyl having from 1-6 carbon atoms;

R_3 is C_{1-6} alkyl or C_{1-6} alkoxy; and

Q_2 is as defined above for Formula I.

Particularly preferred compounds of Formula I-B are those where R_q is ethyl; and

Q_2 is a group of the formula:



wherein R_8 and R_9 together form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

- (a) one or two double bonds;
- (b) one or two oxo;
- (c) one or two O, S or N- R_{10} wherein R_{10} is hydrogen, or C_{1-8} alkyl.

The compounds of Formula I may contain one or more asymmetric centers, so that the compounds can exist in different stereoisomeric forms. All stereoisomeric forms (e.g., optical isomers) and mixtures thereof are encompassed by the invention. The compounds containing asymmetric centers can be, for example, racemates or optically active forms. In these situations, the single enantiomers, i.e., optically active forms, can be obtained by asymmetric synthesis or by resolution of the racemates. Resolution of the racemates can be accomplished, for

example, by conventional methods such as crystallization in the presence of a resolving agent, or chromatography, using, for example a chiral HPLC column.

In addition compounds of Formula I with carbon-carbon
5 double bonds may occur in the Z- and/or E- forms. All such isomeric forms of such compounds are encompassed within the invention. Further, such compounds may be prepared in either pure E or pure Z forms using methods known to those skilled in the art.

10 When any variable (e.g. C₁₋₆ alkyl, C₁₋₈ alkyl, R₁- R₉, Q₁ or Q₂) occurs more than one time in Formula I, its definition on each occurrence is independent of its definition at every other occurrence.

As used herein, the term "alkyl" includes those alkyl
15 groups of a designed number of carbon atoms. Examples of "alkyl" include methyl, ethyl, propyl, isopropyl, butyl, iso-, sec- and tert-butyl, pentyl, hexyl, heptyl, 3-ethylbutyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, norbornyl, and the like. When reference is made herein to C₁₋₆ alkyl or C₁₋₈ alkyl
20 which it may contain one or two double or triple bond it is understood that at least two carbons are present in the alkyl for one double or triple bond, and at least four carbons for two double or triple bonds. "Alkoxy" represents an alkyl group of indicated number of carbon atoms attached through an oxygen
25 bridge, such as methoxy, ethoxy, propoxy and isopropoxy.

By "halogen" herein is meant fluorine, chlorine, bromine, and iodine. Preferred halogens are fluorine and chlorine. Particularly preferred is fluorine.

As used herein, the terms "patient" and "patients" refer to
30 humans as well as other mammals including pets such as dogs and cats and livestock such as cattle and sheep.

By C₁₋₆ alkyl-Ar as used herein is meant an aryl (Ar) group substituted on an alkyl group which is in turn connected to the parent structure. Examples of such groups are benzyl,

phenethyl, 1-naphthylmethyl, 2-pyridylmethyl, and 3-pyrimidin-2-ylpropyl.

This invention also includes methods for using compounds of Formula I to treat diseases or disorders in patients in which
5 mediation by NK-3 receptors is of importance.

The compounds of this invention are ligands for neurokinin receptors, in particular NK-3 receptors, and are useful in the treatment of a wide range of diseases or disorders of the central nervous system (CNS) and periphery in mammals in which
10 modulation of NK-3 receptors is of importance. These include depression, anxiety, panic disorder, obsessive compulsive disorder, psychosis and schizophrenia, neurodegenerative disorders such as dementia, Alzheimer's diseases, Parkinson's disease, Huntington's disease, stress related somatic disorders,
15 reflex sympathetic dystrophy, dysthmic disorders, obesity, eating disorders, drug and alcohol addiction, movement disorders, convulsive disorders such as epilepsy, migraine, headache, multiple sclerosis and other demyelinating diseases, AIDS related neuropathy, chemotherapy-induced neuropathy and
20 neuralgia, diabetic or peripheral neuropathy, neurogenic inflammation, inflammatory pain, neuropathic pain, and other types of chronic or acute pain, Reynaud's disease, vasodilation, vasospasm, angina, asthma, chronic obstructive pulmonary diseases, airway hyperreactivity, cough, allergic
25 rhinitis, bronchospasm, bronchopneumonia, ocular inflammation, inflammatory bowel disease, Crohn's disease, ulcerative colitis, biliary dysfunction, skin disorders and itch, hypersensitivity disorders, atopic dermatitis, contact dermatitis, cutaneous wheal and flare, renal disorders, urinary
30 incontinence, immune system disorders and adverse immunological reactions, fibrositis, osteoarthritis, eosinophilic fascioliasis, and scleroderma.

The present invention also pertains to methods of inhibiting the binding of neurokinin to the NK-3 receptor which

methods involve contacting a compound of the invention with cells expressing NK-3 receptors, wherein the compound is present at a concentration sufficient to inhibit neurokinin binding to cells expressing a cloned human NK-3 receptor *in vitro*. This method includes inhibiting the binding of neurokinin to NK-3 receptors *in vivo*, e.g., in a patient given an amount of a compound of Formula I that would be sufficient to inhibit the binding of neurokinin to NK-3 receptors *in vitro*. The amount of a compound that would be sufficient to inhibit the binding to neurokinin to the NK-3 receptor may be readily determined via an NK-3 receptor binding assay, such as the assay described in Example 35.

The present invention also pertains to methods for altering the signal-transducing activity of NK-3 receptors, said method comprising exposing cells expressing such receptors to an effective amount of a compound of the invention. This method includes altering the signal-transducing activity of NK-3 receptors *in vivo*, e.g., in a patient given an amount of a compound of Formula I that would be sufficient to alter the signal-transducing activity of NK-3 receptors *in vitro*. The amount of a compound that would be sufficient to alter the signal-transducing activity of NK-3 receptors may be determined via an NK-3 receptor signal transduction assay, such as the assay described in Example 36.

The NK-3 antagonist compounds provided by this invention and labeled derivatives thereof are also useful as standards and reagents in determining the ability of a potential pharmaceutical to bind to the NK-3 receptor.

Labeled derivatives the NK-3 antagonist compounds provided by this invention are also useful as radiotracers for positron emission tomography (PET) imaging or for single photon emission computerized tomography (SPECT).

If the compound of the invention is obtained as an acid addition salt, the free base can be obtained by basifying a

solution of the acid salt. Conversely, if the product is a free base, an addition salt, particularly a pharmaceutically acceptable addition salt, may be produced by dissolving the free base in a suitable organic solvent and treating the solution
5 with an acid, in accordance with conventional procedures for preparing acid addition salts from base compounds. Those skilled in the art will recognize various synthetic methodologies that may be employed to prepare non-toxic pharmaceutically acceptable addition salts of the compounds
10 encompassed by Formula I.

Non-toxic pharmaceutical salts include salts of acids such as hydrochloric, phosphoric, diphosphoric, hydrobromic, stearic, sulfuric, sulfinic, formic, fumaric, toluenesulfonic, methanesulfonic, nitric, salicylic, 2-hydroxyethylsulfonic,
15 benzoic, citric, tartaric, lactic, malic, maleic, hydroiodic, alkanolic such as acetic, $\text{HOOC}-(\text{CH}_2)_n-\text{ACOOH}$ where n is 0-4, and the like. Those skilled in the art will recognize a wide variety of non-toxic pharmaceutically acceptable addition salts. Similarly, pharmaceutically acceptable cations include, but are
20 not limited to sodium, potassium, calcium, aluminum, lithium and ammonium.

The present invention also encompasses the prodrugs of the compounds of Formula I. Those skilled in the art will recognize various synthetic methodologies that may be employed to prepare
25 non-toxic pharmaceutically acceptable prodrugs of the compounds encompassed by Formula I. See for example N. Bodor, Drugs of the Future, 1981, 6, 165-182, and H. Bundgaard, Advanced Drug Delivery Reviews, 1989, 3, 39-65.

The compounds of general Formula I may be administered
30 orally, topically, parenterally, by inhalation or spray or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicles. Oral administration in the form of a pill, capsule, elixir, syrup, lozenge, troche, or the like is particularly

preferred. The term parenteral as used herein includes subcutaneous injections, intradermal, intravascular (e.g., intravenous), intramuscular, spinal, intrathecal injection or like injection or infusion techniques. In addition, there is
5 provided a pharmaceutical formulation comprising a compound of general Formula I and a pharmaceutically acceptable carrier. One or more compounds of general Formula I may be present in association with one or more non-toxic pharmaceutically acceptable carriers and/or diluents and/or adjuvants and if
10 desired other active ingredients. The pharmaceutical compositions containing compounds of general Formula I may be in a form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsion, hard or soft capsules, or syrups or elixirs.
15 Compositions intended for oral use may be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavoring agents, coloring agents and
20 preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the active ingredient in admixture with non-toxic pharmaceutically acceptable excipients that are suitable for the manufacture of tablets. These excipients may be for example, inert diluents,
25 such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, for example, corn starch, or alginic acid; binding agents, for example starch, gelatin or acacia, and lubricating agents, for example magnesium stearate, stearic acid or talc.
30 The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed.

Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the
5 active ingredient is mixed with water or an oil medium, for example peanut oil, liquid paraffin or olive oil.

Aqueous suspensions contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients are suspending agents, for
10 example sodium carboxymethylcellulose, methylcellulose, hydropropylmethylcellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents may be a naturally-occurring phosphatide, for example, lecithin, or condensation products of an alkylene oxide
15 with fatty acids, for example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene
20 sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or n-propyl p-hydroxybenzoate, one or more
25 coloring agents, one or more flavoring agents, and one or more sweetening agents, such as sucrose or saccharin.

Oily suspensions may be formulated by suspending the active ingredients in a vegetable oil, for example arachis oil, olive oil, sesame oil or coconut oil, or in a mineral oil such as
30 liquid paraffin. The oily suspensions may contain a thickening agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, and flavoring agents may be added to provide palatable oral preparations.

These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

Pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, for example olive oil or arachis oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying agents may be naturally-occurring gums, for example gum acacia or gum tragacanth, naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol, anhydrides, for example sorbitan monoleate, and condensation products of the said partial esters with ethylene oxide, for example polyoxyethylene sorbitan monoleate. The emulsions may also contain sweetening and flavoring agents.

Syrups and elixirs may be formulated with sweetening agents, for example glycerol, propylene glycol, sorbitol or sucrose. Such formulations may also contain a demulcent, a preservative and flavoring and coloring agents. The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to the known art using those suitable dispersing or wetting agents and suspending agents which have been mentioned above. The sterile injectable preparation may also be sterile injectable solution or suspension in a non-toxic parentally acceptable diluent or solvent, for example as a solution in 1,3-butanediol. Among the

acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed
5 oil may be employed including synthetic mono-or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

The compounds of general Formula I may also be administered in the form of suppositories, e.g., for rectal administration of
10 the drug. These compositions can be prepared by mixing the drug with a suitable non-irritating excipient that is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

15 Compounds of general Formula I may be administered parenterally in a sterile medium. The drug, depending on the vehicle and concentration used, can either be suspended or dissolved in the vehicle. Advantageously, adjuvants such as local anesthetics, preservatives and buffering agents can be
20 dissolved in the vehicle.

Dosage levels of the order of from about 0.1 mg to about 140 mg per kilogram of body weight per day are useful in the treatment of the above-indicated conditions (about 0.5 mg to about 7 g per patient per day). The amount of active ingredient
25 that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. Dosage unit forms will generally contain between from about 1 mg to about 500 mg of an active ingredient.

30 Frequency of dosage may also vary depending on the compound used and the particular disease treated. However, for treatment of most disorders, a dosage regimen of 4 times daily or less is preferred. For the treatment of schizophrenia, depression, or

obesity a dosage regimen of 1 or 2 times daily is particularly preferred.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, and rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

Preferred compounds of the invention have certain pharmacological properties. Such properties include, but are not limited to oral bioavailability, low toxicity, low serum protein binding and desirable *in vitro* and *in vivo* half-lives. Penetration of the blood brain barrier for compounds used to treat CNS disorders is necessary, while low brain levels of compounds used to treat peripheral disorders are generally preferred.

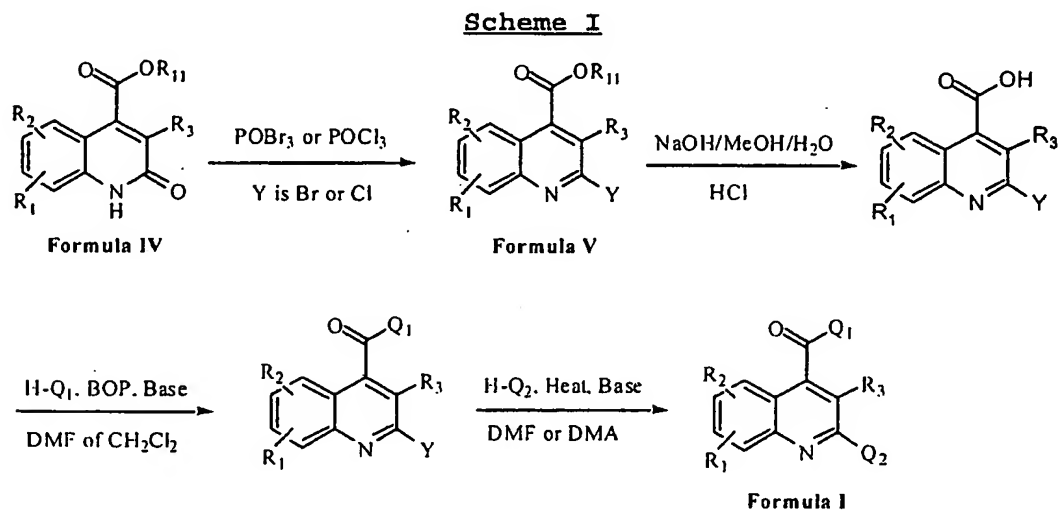
Assays may be used to predict these desirable pharmacological properties. Assays used to predict bioavailability include transport across human intestinal cell monolayers, including Caco-2 cell monolayers. Toxicity to cultured hepatocytes may be used to predict compound toxicity. Penetration of the blood brain barrier of a compound in humans may be predicted from the brain levels of the compound in laboratory animals given the compound intravenously.

Serum protein binding may be predicted from albumin binding assays. Such assays are described in a review by Oravcová, et al. (Journal of Chromatography B (1996) volume 677, pages 1-27).

Compound half-life is inversely proportional to the frequency of dosage of a compound. *In vitro* half-lives of compounds may be predicted from assays of microsomal half-life as described by Kuhnz and Gieschen (Drug Metabolism and Disposition, (1998) volume 26, pages 1120-1127).

The present invention also pertains to packaged pharmaceutical compositions for treating disorders responsive to NK-3 receptor modulation, e.g., treatment of schizophrenia, depression, or chronic pulmonary obstructive disorder by NK-3 receptor modulation. The packaged pharmaceutical compositions include a container holding a therapeutically effective amount of at least one NK-3 receptor modulator as described supra and instructions (e.g., labeling) indicating the contained NK-3 receptor ligand is to be used for treating a disorder responsive to NK-3 receptor modulation in the patient.

Compounds of Formula I wherein X is O are prepared by the method described in Scheme I.



15

In Scheme I, R_1 , R_2 , R_3 , Q_1 and Q_2 are as defined above for Formula I, and R_{11} is methyl, ethyl or benzyl. BOP is benzotriazol-1-yloxytris(dimethylamino)-phosphoniumhexafluoro-phosphate. Base is triethylamine, diisopropylethylamine, *N*-methylmorpholine or other suitable organic base. DMF is dimethylformide. DMA is dimethylacetamide. Heat as used herein means elevated temperature, such as, for example, 40 to 250°C.

Compounds of the invention where X is S or N-CN can be prepared using similar reactions. Alternatively, the compounds

of Formula I where X is O can be converted to the corresponding compounds where X is S or N-CN.

Compounds of Formula IV are prepared using literature procedures several publications including Jacobs, T. L. et al.;
5 Org. Synth. Coll. 1955, Vol. 3, 456-58 and Lyle R. E. et al. J. Org. Chem. 1972, 37, 3967-68 if they are not commercially available. Compounds of Formula V are prepared by a method described in Batt D. G. et al. Bioorg. Med. Chem. Lett. 1998, 8, 1745-50. Those skilled in the art will recognize that in
10 certain instances it will be necessary to utilize compounds of Formula IV bearing protecting groups and that these groups can be removed in a subsequent reaction to yield compounds of Formula I as described in "Protective Groups in Organic Synthesis", 2nd Ed., Greene, T. W. and related publications.

15 In some situations, the replacement of Y with Q₂ or a protected form of Q₂ may be facilitated by use of a palladium catalyst as described in Buchwald, S. L. et al in J. Org. Chem. 2000, 65, 1144-57 and 1158-74. Typical conditions for this reaction include but are not limited to treatment of the 2-
20 chloro or 2-bromoquinoline derivative with Q₂H, sodium tert-butoxide, catalytic tris(dibenzylideneacetone)-dipalladium(0) and catalytic BINAP ([1,1'-bisnaphthalene]-2,2'-diylbis(diphenylphosphine)) in toluene at 50-120 °C. These reaction conditions are often useful for efficient conversion of
25 Y to Q₂ when R₃ is not hydrogen.

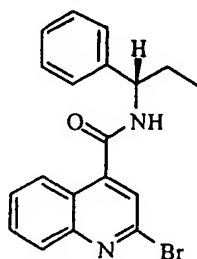
Those having skill in the art will recognize that the starting materials, solvents, and other reaction conditions may be varied and additional steps employed to produce compounds encompassed by the present invention. In some cases protection
30 of certain reactive functionalities may be necessary to achieve some of the above transformations. In general the need for such protecting groups will be apparent to those skilled in the art of organic synthesis as well as the conditions necessary to attach and remove such groups.

The invention is illustrated further by the following examples, which are not to be construed as limiting the invention in scope or spirit to the specific procedures described in them.

5 The disclosures of all articles and references mentioned in this document, including patents, are incorporated herein by reference.

Example 1

10 (S)-N-(α -Ethylphenyl)-2-bromoquinoline-4-carboxamide

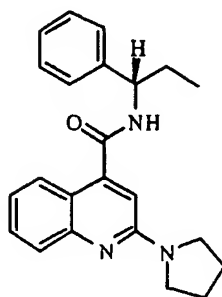


A mixture of methyl 2-bromoquinolinecarboxylate (7.2 g) and NaOH (1.1 g) in 15 mL of methanol and 15 mL of water is stirred at 50 °C until TLC indicates the disappearance of starting
15 material. After cooling, the methanol is evaporated and the mixture is treated with 28 mL of 1N HCl, the solid is collected by filtration and dried to afford 6.1 g of 2-bromoquinoline-4-carboxylic acid.

To a solution of 2-bromoquinoline-4-carboxylic acid (2.6 g)
20 in 25 mL of DMF containing is added BOP (5.0 g), followed by (S)-1-phenylpropylamine (2.8 g). The reaction mixture is stirred overnight at room temperature. The mixture is diluted with 20 mL of CH₂Cl₂, then washed with aqueous NaHCO₃, aqueous citric acid and brine. After drying over Na₂SO₄, the solvent is evaporated to
25 afford the corresponding compound is obtained as a white solid.

Example 2

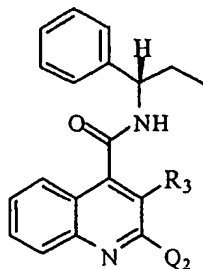
(S)-N-(α -Ethylphenyl)-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide



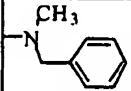
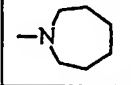
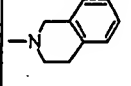
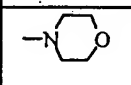
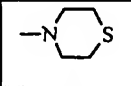
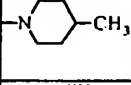
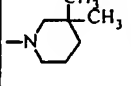
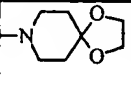
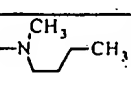
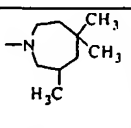
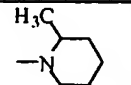
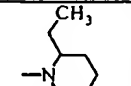
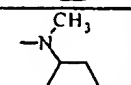
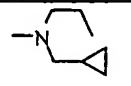
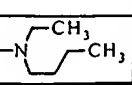
A solution of (*S*)-*N*-(α -Ethylphenyl)-2-bromoquinoline-4-carboxamide (500 mg), pyrroline (1 mL) and triethylamine (1 mL) in 5 mL of DMF in a sealed tube is heated with stirring for 8 hours. The reaction mixture is poured into ice-water. The product is collected by filtration, washed with water and dried. The corresponding compound is obtained as a tan solid.

The following examples listed in Table 1 are prepared using procedures analogous to those set forth above. MS data are given as $[MH]^+$. LC-MS data are given as HPLC retention times and $[MH]^+$. The HPLC retention times shown in Table 1 are obtained by essentially according to the method described in Example 1.

Table 1



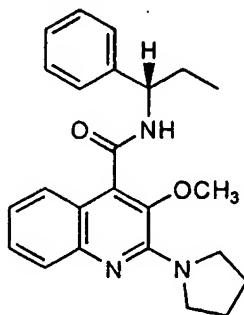
Example	R ₃	-Q ₂	Compound Name	HPLC retention time (min)	$[MH]^+$
3	H		(<i>S</i>)- <i>N</i> -(α -ethylphenyl)-2-(piperidin-1-yl)-quinoline-4-carboxamide	NA*	374
4	H		(<i>S</i>)- <i>N</i> -(α -ethylphenyl)-2-(1,2,5,6-	2.17	372

			tetrahydropyridin-1-yl)-quinoline-4-carboxamide		
5	H		(S)-N-(α -ethylphenyl)-2-(N,N-methylbenzylamino)-quinoline-4-carboxamide	2.30	410
6	H		(S)-N-(α -ethylphenyl)-2-(homopiperidin-1-yl)-quinoline-4-carboxamide	2.23	388
7	H		(S)-N-(α -ethylphenyl)-2-(1,2,3,4-tetrahydroisoquinolin-2-yl)-quinoline-4-carboxamide	2.30	422
8	H		(S)-N-(α -ethylphenyl)-2-(morpholin-4-yl)-quinoline-4-carboxamide	2.09	376
9	H		(S)-N-(α -ethylphenyl)-2-(thiomorpholin-4-yl)-quinoline-4-carboxamide	2.17	392
10	H		(S)-N-(α -ethylphenyl)-2-(4-methylpiperidin-1-yl)-quinoline-4-carboxamide	2.26	389
11	H		(S)-N-(α -ethylphenyl)-2-(3,3-dimethylpiperidin-1-yl)-quinoline-4-carboxamide	2.31	402
12	H		(S)-N-(α -ethylphenyl)-2-(aza-1,4-dioxaspiro[4.5]decanyl-8-yl)-quinoline-4-carboxamide	2.17	432
13	H		(S)-N-(α -ethylphenyl)-2-(N-methyl-N-n-butylamino)-quinoline-4-carboxamide	2.27	376
14	H		(S)-N-(α -Ethylphenyl)-2-(3,5,5-trimethylhomopiperidin-1-yl)-quinoline-4-carboxamide	2.47	430
15	H		(S)-N-(α -Ethylphenyl)-2-(2-methylpiperidin-1-yl)-quinoline-4-carboxamide	2.22	388
16	H		(S)-N-(α -ethylphenyl)-2-(2-ethylpiperidin-1-yl)-quinoline-4-carboxamide	2.27	402
17	H		(S)-N-(α -ethylphenyl)-2-(N-methyl-N-cyclohexylamino)-quinoline-4-carboxamide	2.27	402
18	H		(S)-N-(α -ethylphenyl)-2-(N-n-propyl-N-cyclopropylmethylamino)-quinoline-4-carboxamide	2.29	402
19	H		(S)-N-(α -ethylphenyl)-2-(N-ethyl-N-n-butylamino)-quinoline-4-carboxamide	2.30	390

			butylamino)-quinoline-4-carboxamide		
20	H		(S)-N-(α-ethylphenyl)-2-(4-methylpiperazin-1-yl)-quinoline-4-carboxamide	2.09	389
21	H		(S)-N-(α-ethylphenyl)-2-(3-hydroxypyrrolidin-1-yl)-quinoline-4-carboxamide	2.07	376
22	H		(S)-N-(α-ethylphenyl)-2-[(S)-2-methoxymethylpyrrolidin-1-yl]-quinoline-4-carboxamide	2.18	404
23	H		(S)-N-(α-ethylphenyl)-2-[(S)-2-hydroxymethylpyrrolidin-1-yl]-quinoline-4-carboxamide	2.09	390
24	H		(S)-N-(α-ethylphenyl)-2-(2-carboxypyrrolidin-1-yl)-quinoline-4-carboxamide	2.12	404
25	H		(S)-N-(α-ethylphenyl)-2-(3-hydroxymethylpiperidin-1-yl)-quinoline-4-carboxamide	2.13	404
26	H		(S)-N-(α-ethylphenyl)-2-(2-hydroxymethylpiperidin-1-yl)-quinoline-4-carboxamide	NA	404
27	H		(S)-N-(α-ethylphenyl)-2-(3-hydroxypiperidin-1-yl)-quinoline-4-carboxamide	2.09	390
28	H		(S)-N-(α-ethylphenyl)-2-[2-(1-hydroxyethyl)piperidin-1-yl]-quinoline-4-carboxamide	NA	418
29	H		(S)-N-(α-Ethylphenyl)-2-(N-α-methylbenzylamino)-quinoline-4-carboxamide	NA	410
30	H		(S)-N-(α-Ethylphenyl)-2-(azabicyclo[3.2.2]nonan-3-yl)-quinoline-4-carboxamide.	2.30	414
31	CH ₃		(S)-N-(α-Ethylphenyl)-3-methyl-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide	NA	374
32	CH ₃		(S)-N-(α-Ethylphenyl)-3-methyl-2-(piperidin-1-yl)-quinoline-4-carboxamide	NA	388
33	OCH ₃		(S)-N-(α-Ethylphenyl)-3-methoxy-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide	NA	NA
34	OCH ₃		(S)-N-(α-Ethylphenyl)-3-methoxy-2-(piperidin-1-yl)-quinoline-4-carboxamide	NA	NA

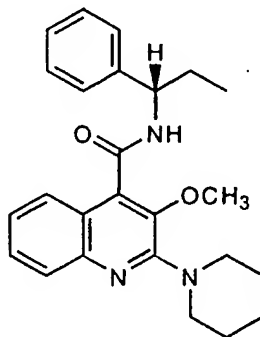
NA: not determined

The compound of Example 33 can be prepared according to the
5 following procedure.



A solution of (*S*)-*N*-(α-Ethylphenyl)-2-bromo-3-methoxyquinoline-
4-carboxamide (500 mg), pyrrolidine (106 mg), sodium *tert*-butoxide
(1.75 mmol), Pd₂(dba)₃ (0.003 mmol) and BINAP (0.0094 mmol) in
10 toluene (2.5 ml) under argon is immersed in an oil bath at 80 °C
with magnetic stirring. After complete consumption of starting
bromide, the reaction mixture is allowed to cool to ambient
temperature, diluted with ethyl acetate (10 ml), filtered,
concentrated and purified by chromatography on silica gel to
15 yield the desired product.

The compound of Example 34 can be prepared according to the
following procedure.



A solution of (*S*)-*N*-(α-Ethylphenyl)-2-bromo-3-methoxyquinoline-
4-carboxamide (500 mg), piperidine (128 mg), sodium *tert*-butoxide
(1.75 mmol), Pd₂(dba)₃ (0.003 mmol) and BINAP (0.0094 mmol) in
20 toluene (2.5 ml) under argon is immersed in an oil bath at 80 °C
with magnetic stirring. After complete consumption of starting
25 bromide, the reaction mixture is allowed to cool to ambient

temperature, diluted with ethyl acetate (10 ml), filtered, concentrated and purified by chromatography on silica gel to yield the desired product.

5

Example 35

NK-3 Receptor Binding Assay

The following assay is a standard assay of NK3 receptor binding, which is used to determine the NK-3 receptor binding
10 affinity of compounds.

Assays are performed as described in Krause et al (Proc. Natl. Acad. Sci. USA 94: 310-315, 1997). The NK-3 receptor complementary DNA was cloned from human hypothalamic RNA using standard procedures. The receptor cDNA was inserted into the
15 expression vector pM² to transfect the mammalian Chinese hamster ovary cell line, and a stably expressing clonal cell line was isolated, characterized and used for the current experiments. Cells are grown in minimal essential medium alpha containing 10% fetal bovine serum and 0.8 mg G418 per ml. Cells were liberated
20 from cell culture plates with No-zyne (PBS base, JRH Biosciences), and harvested by low speed centrifugation. The cell pellet was homogenized in TBS (0.05 M TrisHCl, 120 mM NaCl, pH 7.4) with a Polytron homogenizer at setting 5 for 20 seconds, and total cellular membranes were isolated by centrifugation at
25 47,500 x g for 10 minutes. The membrane pellet was resuspended by homogenization with the Polytron as above, and the membranes were isolated by centrifugation at 47,500 x g for 10 minutes. This final membrane pellet was resuspended in TBS at a protein concentration of 350 µg/ml.

30 Receptor binding assays contain a total volume of 200 µl containing 50 µg membrane protein, 0.15 nM ¹²⁵I-methylPhe⁷-neurokinin B, drug or blocker in TBS containing 1.0 mg/ml bovine serum albumen, 0.2 mg/ml bacitracin, 20 µg/ml leupeptin and 20 µg/ml chymostatin. Incubations are carried out for 2 hours at

4°, and the membrane proteins are harvested by passing the incubation mixture by rapid filtration over presoaked GF/B filters to separate bound from free ligand. The filters are presoaked in TBS containing 2% BSA and 0.1% Tween 20. After
5 filtration of the incubation mixture, filters are rinsed 4 times with ice-cold TBS containing 0.01% sodium dodecyl sulfate and counted in a β -plate scintillation counter. One μ M methylPhe⁷-neurokinin B is added to some tubes to determine nonspecific binding. Data are collected in duplicate determinations,
10 averaged, and the percent inhibition of total specific binding is calculated. The total specific binding is the total binding minus the nonspecific binding. In many cases, the concentration of unlabeled drug is varied and total displacement curves of binding are carried out. Data are converted to a form for the
15 calculation of IC₅₀ and Hill coefficient (nH). Compounds of Formula I display IC₅₀ values less than 5 μ M. Preferred compounds of Formula I display IC₅₀ values of less than 1 μ M, more preferred compounds of Formula I display IC₅₀ values of less than 1 μ M, still more preferred compounds of Formula I display IC₅₀
20 values of less than 100 nM, and particularly preferred compounds of Formula I display IC₅₀ values of less than 10 nM.

Example 36

Assay For NK-3 Functional Activity

25 Calcium Mobilization Assays: The human NK-3 bearing Chinese hamster ovary cells are grown in minimal essential media supplemented with 250 ug/ml G418, 10% fetal bovine serum and 25 mM Hepes, pH=7.4. Forty eight hours prior to the day of assay, the cells are plated in fresh media that does not contain the
30 G418. On the day of assay, cells grown to 70-90% confluency in 96-well plates are washed with Krebs-Ringer buffer (25 mM HEPES, 5 mM KCl, 0.96 mM NaH₂PO₄, 1 mM MgSO₄, 2 mM CaCl₂, 5 mM glucose, 1 mM probenecid, pH 7.4) and are then incubated for 1-2 hours in

the above buffer supplemented with Fluo3-AM (2.5 to 10 µg/ml; Teflabs) at 37 degrees C in an environment containing 5% CO₂. The wells are then washed twice with Krebs Ringers HEPES buffer. Agonist-induced (methylPhe7-neurokinin B) calcium mobilization is monitored using a FLIPR (Molecular Devices) instrument. The agonist is added to the cells and fluorescence responses are continuously recorded for up to 5min. For the examination of antagonist drug candidates, compounds are preincubated with the cells for up to 30 min. prior to administration of the methylPhe7-neurokinin B agonist usually at a concentration that brings about a 50% maximal activity. Responses are recorded for times up to 5 min. Kaleidagraph software (Synergy Software, Reading, PA) is utilized to fit the data to the equation $y = a * (1 / (1 + (b/x)^c))$ to determine the EC₅₀ value or IC₅₀ value for the response. In this equation, y is the maximum fluorescence signal, x is the concentration of the agonist or antagonist, a is the E_{max}, b corresponds to the EC₅₀ or IC₅₀ value, and, finally, c is the Hill coefficient.

20 Example 37

Preparation of radiolabeled probe compounds of the invention

The compounds of the invention are prepared as radiolabeled probes by carrying out their synthesis using precursors comprising at least one atom that is a radioisotope. The radioisotope is preferably selected from of at least one of carbon (preferably ¹⁴C), hydrogen (preferably ³H), sulfur (preferably ³⁵S), or iodine (preferably ¹²⁵I). A commercial laboratory specializing in custom synthesis of radiolabeled probe compounds conveniently carries out synthesis of such radiolabeled probes. Such laboratories include Amersham Corporation, Arlington Heights, IL; Cambridge Isotope Laboratories, Inc. Andover, MA; SRI International, Menlo Park, CA; Wizard Laboratories, West Sacramento, CA; ChemSyn

Laboratories, Lexena, KS; American Radiolabeled Chemicals, Inc., St. Louis, MO; and Moravsek Biochemicals Inc., Brea, CA.

5 Tritium-labeled probe compounds are also conveniently prepared catalytically via platinum-catalyzed exchange in tritiated acetic acid, acid-catalyzed exchange in tritiated trifluoroacetic acid, or heterogeneous-catalyzed exchange with tritium gas. Such preparations are also conveniently carried out as a custom radiolabeling by any of the suppliers listed in the preceding paragraph using the compound of the invention as
10 substrate. In addition, certain precursors may be subjected to tritium-halogen exchange with tritium gas, tritium gas reduction of unsaturated bonds, or reduction using sodium borotritide, as appropriate.

15 Example 38

Use of compounds of the invention as probes for detecting NK-3 receptors in cultured cells and tissue samples (e.g., tissue sections)

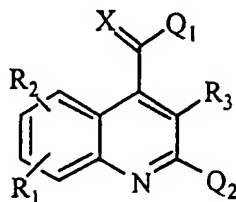
20 Receptor autoradiography (receptor mapping) of NK-3 receptors in cultured cells or tissue samples is carried out in vitro as described by Kuhar in sections 8.1.1 to 8.1.9 of Current Protocols in Pharmacology (1998) John Wiley & Sons, New York, using radiolabeled compounds of the invention prepared as described in the preceding Example.

25 The invention and the manner and process of making and using it, are now described in such full, clear, concise and exact terms as to enable any person skilled in the art to which it pertains, to make and use the same. It is to be understood that the foregoing describes preferred embodiments of the
30 present invention and that modifications may be made therein without departing from the spirit or scope of the present invention as set forth in the claims. To particularly point out

and distinctly claim the subject matter regarded as invention,
the following claims conclude this specification.

WHAT IS CLAIMED:

1. A compound of the formula:



5 or pharmaceutically acceptable salts or pharmaceutically acceptable solvates thereof, wherein:

R₁ is selected from:

hydrogen, halogen, hydroxy, C₁₋₆ alkyl, -O(C₁₋₆ alkyl), -NO₂,
 10 -CN, -SO₂NH₂, -SO₂NH(C₁₋₆ alkyl), -SO₂N(C₁₋₆ alkyl)(C₁₋₆ alkyl),
 amino, -NH(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)(C₁₋₆ alkyl), -N(C₁₋₆
 alkyl)CO(C₁₋₆ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)CO₂(C₁₋₆
 alkyl), -NHSO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)SO₂(C₁₋₆ alkyl),
 -SO₂NHCO(C₁₋₆ alkyl), -CONHSO₂(C₁₋₆ alkyl), -CON(C₁₋₆ alkyl)
 15 (C₁₋₆ alkyl), -CO₂(C₁₋₆ alkyl), -S(C₁₋₆ alkyl), -SO(C₁₋₆ alkyl),
 or -SO₂(C₁₋₆ alkyl),

wherein said C₁₋₆ alkyl may be straight, branched or cyclic,
 may contain one or two double or triple bonds,
 unsubstituted or substituted with one or more substituents
 20 selected from: hydroxy, oxo, halogen, amino, or C₁₋₃ alkoxy;

R₂ and R₃ are independently selected from the group consisting
 of:

hydrogen, halogen, hydroxy, C₁₋₈ alkyl, -O(C₁₋₈ alkyl), -NO₂,
 -CN, -SO₂NH₂, -SO₂NH(C₁₋₈ alkyl), -SO₂N(C₁₋₈ alkyl)(C₁₋₈ alkyl),
 25 amino, -NH(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)(C₁₋₈ alkyl), -N(C₁₋₈
 alkyl)CO(C₁₋₈ alkyl), -NHCO₂(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)CO₂(C₁₋₈
 alkyl), -NHSO₂(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)SO₂(C₁₋₈ alkyl), -
 SO₂NHCO(C₁₋₈ alkyl), -CONHSO₂(C₁₋₈ alkyl), -CON(C₁₋₈ alkyl)(C₁₋₈
 alkyl), -CO₂(C₁₋₈ alkyl), -S(C₁₋₈ alkyl), -SO(C₁₋₈ alkyl), -
 30 SO₂(C₁₋₈ alkyl), and Ar,

wherein Ar is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, benzoimidazolyl, furanyl, benzofuranyl, thiazolyl, benzothiazolyl, isothiazolyl, benzisothiazolyl, triazolyl, tetrazolyl, pyrazolyl, or benzopyrazolyl, each of which is unsubstituted or substituted with one or more substituents selected from:

halogen, hydroxy, C₁₋₆ alkyl, -O(C₁₋₆ alkyl), -NO₂, -CN, -SO₂NH₂, -SO₂NH(C₁₋₆ alkyl), -SO₂N(C₁₋₆ alkyl)(C₁₋₆ alkyl), amino, -NH(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)CO(C₁₋₆ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)CO₂(C₁₋₆ alkyl), -NHSO₂(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)SO₂(C₁₋₆ alkyl), -SO₂NHCO(C₁₋₆ alkyl), -CONHSO₂(C₁₋₆ alkyl), -CON(C₁₋₆ alkyl)(C₁₋₆ alkyl), -CO₂(C₁₋₆ alkyl), -S(C₁₋₆ alkyl), -SO(C₁₋₆ alkyl), or -SO₂(C₁₋₆ alkyl), wherein C₁₋₆ alkyl, is defined as above,

wherein said C₁₋₈ alkyl and said C₁₋₆ alkyl may be straight, branched or cyclic, may contain one or two double or triple bonds, unsubstituted or substituted with one or more of the substituents selected from:

- (i) hydroxy,
- (ii) oxo,
- (iii) halogen,
- (iv) Ar, wherein Ar is as defined above,
- (v) -NR₄R₅, wherein R₄ and R₅ are independently selected from:

- (A) hydrogen,
- (B) C₁₋₆ alkyl, wherein C₁₋₆ alkyl is as defined above,
- (C) Ar, where in Ar is as defined above,
- (D) C₁₋₆ alkyl-Ar, wherein Ar is as defined above and Ar is attached to any position of the C₁₋₆ alkyl group at any position of Ar,

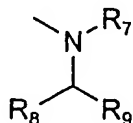
and the groups R_4 and R_5 may be joined together to form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

- (a) one or two double bonds,
- 5 (b) one or two oxo,
- (c) one or two O, S or N- R_6 wherein R_6 is hydrogen, C_{1-6} alkyl, C_{1-6} alkyl-Ar, wherein C_{1-6} alkyl and Ar are defined as above and Ar is attached to C_{1-6} alkyl at any position,
- 10 (d) one or two R_1 ,
- (vi) $-OR_4$, wherein R_4 is as defined above,
- (vii) $-CONR_4R_5$, wherein R_4 and R_5 are as defined above,
- (viii) $-SO_2NR_4R_5$, wherein R_4 and R_5 are as defined above,
- (ix) $-NR_4COR_5$, wherein R_4 and R_5 are as defined above,

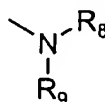
15

X is O, S or N-CN;

Q_1 and Q_2 are independently selected from formulas II and III:



II



III

20

wherein R_7 is hydrogen or C_{1-8} alkyl, wherein C_{1-8} alkyl is as defined above,

wherein R_8 and R_9 are independently selected from:

- 25 (i) hydrogen,
- (ii) Ar' , wherein Ar' is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, benzoimidazolyl, furanyl, benzofuranyl, thiazolyl, benzothiazolyl, isothiazolyl,
- 30 benzisothiazolyl, triazolyl, tetrazolyl, pyrazolyl, or benzopyrazolyl, each of which is unsubstituted or substituted with one or more substituents selected from:

- halogen, hydroxy, C₁₋₈ alkyl, -O(C₁₋₈ alkyl), -NO₂, -CN, -SO₂NH₂, -SO₂NH(C₁₋₈ alkyl), -SO₂N(C₁₋₈ alkyl)(C₁₋₈ alkyl), amino, -NH(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)(C₁₋₈ alkyl), -N(C₁₋₈ alkyl)CO(C₁₋₈ alkyl), -NHCO₂(C₁₋₆ alkyl), -N(C₁₋₈ alkyl)CO₂(C₁₋₈ alkyl), -NHCO₂(C₁₋₈ alkyl), -N(C₁₋₆ alkyl)SO₂(C₁₋₈ alkyl), -SO₂NHCO(C₁₋₈ alkyl), -CONHSO₂(C₁₋₈ alkyl), -CON(C₁₋₈ alkyl)(C₁₋₈ alkyl), -CO₂(C₁₋₈ alkyl), -S(C₁₋₈ alkyl), -SO(C₁₋₈ alkyl), and -SO₂(C₁₋₈ alkyl), wherein C₁₋₈ alkyl, is defined as above,
- (iii) C₁₋₈ alkyl, wherein said C₁₋₈ alkyl is as defined above,
- (iv) -COR₄, wherein R₄ is as defined above,
- (v) -CONR₄R₅, wherein R₄ and R₅ are as defined above,
- (vi) -SO₂NR₄R₅, wherein R₄ and R₅ are as defined above
- or the groups R₄ and R₅ may be joined together to form a 4- to 8-membered monocyclic or bicyclic ring of which may contain:
- (a) one or two double bonds,
- (b) one or two oxo,
- (c) one or two O, S or N-R₁₀ wherein R₁₀ is hydrogen, C₁₋₈ alkyl, C₁₋₈ alkyl-Ar', wherein C₁₋₈ alkyl and Ar' are as defined above and Ar' is attached to any position of the C₁₋₈ alkyl group at any position of Ar';
- (d) one or two R₁.

25

2. A compound according to claim 1, which is (S)-N-(α-ethylphenyl)-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide.

3. A compound according to claim 1, which is (S)-N-(α-ethylphenyl)-2-(piperidin-1-yl)-quinoline-4-carboxamide.

30

4. A compound according to claim 1, which is (S)-N-(α-ethylphenyl)-2-(1,2,5,6-tetrahydropyridin-1-yl)-quinoline-4-carboxamide.

5. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(N,N-methylbenzylamino)-quinoline-4-carboxamide.

5 6. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(homopiperidin-1-yl)-quinoline-4-carboxamide.

7. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(1,2,3,4-tetrahydroisoquinolin-2-yl)-quinoline-4-
10 carboxamide.

8. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(morpholin-4-yl)-quinoline-4-carboxamide.

15 9. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(thiomorpholin-4-yl)-quinoline-4-carboxamide.

10. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(4-methylpiperidin-1-yl)-quinoline-4-carboxamide.
20

11. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(3,3-dimethylpiperidin-1-yl)-quinoline-4-carboxamide.

25 12. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(aza-1,4-dioxaspiro[4.5]decanyl-8-yl)-quinoline-4-carboxamide.

13. A compound according to claim 1, which is (S)-N-(α -
30 ethylphenyl)-2-(N-methyl-N-n-butylamino)-quinoline-4-carboxamide.

14. A compound according to claim 1, which is (S)-N-(α -Ethylphenyl)-2-(3,,5,5-trimethylhomopiperidin-1-yl)-quinoline-4-carboxamide.

5 15. A compound according to claim 1, which is (S)-N-(α -Ethylphenyl)-2-(2-methylpiperidin-1-yl)-quinoline-4-carboxamide.

16. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(2-ethylpiperidin-1-yl)-quinoline-4-carboxamide.

10

17. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(N-methyl-N-cyclohexylamino)-quinoline-4-carboxamide.

15 18. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(N-n-propyl-N-cyclopropylmethylamino)-quinoline-4-carboxamide.

19. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(N-ethyl-N-n-butylamino)-quinoline-4-carboxamide.

20

20. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(4-methylpiperazin-1-yl)-quinoline-4-carboxamide.

25 21. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(3-hydroxypyrrolidin-1-yl)-quinoline-4-carboxamide.

22. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-[(S)-2-methoxymethylpyrrolidin-1-yl]-quinoline-4-carboxamide.

30

23. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-[(S)-2-hydroxymethylpyrrolidin-1-yl]-quinoline-4-carboxamide.

5 24. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(2-carboxypyrrolidin-1-yl)-quinoline-4-carboxamide.

10 25. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(3-hydroxymethylpiperidin-1-yl)-quinoline-4-carboxamide.

15 26. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(2-hydroxymethylpiperidin-1-yl)-quinoline-4-carboxamide.

20 27. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(3-hydroxypiperidin-1-yl)-quinoline-4-carboxamide.

28. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-[2-(2-hydroxyethyl)piperidin-1-yl]-quinoline-4-carboxamide.

25 29. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(N- α -methylbenzylamino)-quinoline-4-carboxamide.

30 30. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-2-(azabicyclo[3.2.2]nonan-3-yl)-quinoline-4-carboxamide.

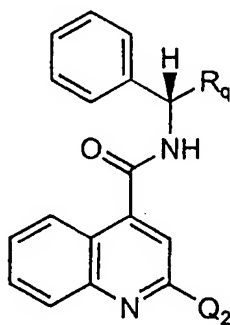
31. A compound according to claim 1, which is (S)-N-(α -Ethylphenyl)-3-methyl-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide.

5 32. A compound according to claim 1, which is (S)-N-(α -Ethylphenyl)-3-methyl-2-(piperidin-1-yl)-quinoline-4-carboxamide.

33. A compound according to claim 1, which is (S)-N-(α -Ethylphenyl)-3-methoxy-2-(pyrrolidin-1-yl)-quinoline-4-carboxamide.

34. A compound according to claim 1, which is (S)-N-(α -ethylphenyl)-3-methoxy-2-(piperidin-1-yl)-quinoline-4-carboxamide.

35. A compound according to claim 1, of the formula:



20 where R_q is straight or branched chain alkyl having from 1-6 carbon atoms; and
 Q_2 is as defined in Claim 1.

25 36. A compound according to claim 35, wherein:

R_q is ethyl; and

R_8 and R_9 are independently selected from:

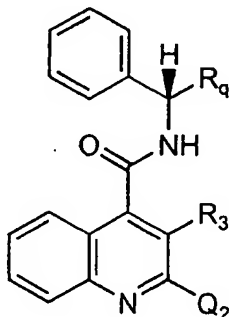
hydrogen;

C₁₋₈ alkyl, wherein said C₁₋₈ alkyl is as defined above;
or

R₈ and R₉ together form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

- 5 (a) one or two double bonds;
- (b) one or two oxo;
- (c) one or two O, S or N-R₁₀ wherein R₁₀ is hydrogen, C₁₋₈ alkyl, C₁₋₈ alkyl-Ar', wherein C₁₋₈ alkyl and Ar' are defined as above and Ar' may be attached to C₁₋₈ alkyl at
- 10 any position; or
- (d) one or two R₁ groups.

37. A compound according to Claim 1, of the formula:



15 wherein:

R_q is straight or branched chain alkyl having from 1-6 carbon atoms;

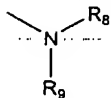
20 R₃ is C₁₋₆ alkyl or C₁₋₆ alkoxy; and

Q₂ is as defined in Claim 1.

38. A compound according to Claim 37, wherein:

R_q is ethyl; and

25 Q₂ is a group of the formula:



wherein R₈ and R₉ together form a 4- to 8-membered monocyclic or bicyclic ring which may contain:

(a) one or two double bonds;

(b) one or two oxo;

5 (c) one or two O, S or N-R₁₀ wherein R₁₀ is hydrogen, or C₁₋₈ alkyl.

39. All stereoisomeric forms of compounds described in claims 1 through 30 and mixtures thereof.

10

40. A compound according to Claim 1, for use in
15 therapeutic treatment of a disease or disorder associated with pathogenic neurokinin-3 receptor activation.

41. A compound according to Claim 1, for use in the treatment of immune disorders, migraine, biliary dysfunction, and dermatitis in patients.

20

42. A compound according to Claim 1, for use in the treatment of Parkinson's disease, Alzheimer's disease, neurodegenerative diseases, neuropathy, or movement disorders.

25 43. A compound according to Claim 1, for use in the treatment of anxiety, depression, sleep disorders, drug addiction, schizophrenia, obesity, chronic pulmonary obstructive disorder respiratory diseases, inflammatory diseases, or pain.

30 44. The use of a compound according to Claim 1 for the manufacture of a medicament for the treatment of a disease or disorder associated with pathogenic neurokinin-3 receptor activation.

45. The use of a compound according to Claim 1 for the manufacture of a medicament for the treatment of anxiety, depression, sleep disorders, drug addiction, schizophrenia, obesity, chronic pulmonary obstructive disorder or pain.

5

46. A pharmaceutical composition comprising a compound according to Claim 1 combined with at least one pharmaceutically acceptable carrier or excipient.

10

47. A method for the treatment of a disease or disorder associated with pathogenic neurokinin-3 receptor activation, said method comprising administering to a patient in need of such treatment an effective amount of a compound of claim 1.

15

48. A method according to Claim 47 wherein the disorder associated with pathogenic neurokinin-3 receptor is anxiety, depression, schizophrenia, obesity, chronic pulmonary obstructive disorder, or pain.

20

49. A method according to Claim 47 wherein the disorder associated with pathogenic neurokinin-3 receptor is an immune disorders, migraine, biliary disfunction, or dermatitis.

25

50. A method according to Claim 47 wherein the disorder associated with pathogenic neurokinin-3 receptor is Parkinson's disease, Alzheimer's disease, a neurodegenerative disease, neuropathy, or a movement disorder.

30

51. A method for localizing neurokinin-3 receptors in a tissue sample comprising:
contacting with the sample a detectably-labeled compound of claim 1 under conditions that permit binding of the compound to neurokinin 3 receptors, washing the sample to remove unbound compound, and detecting the bound compound.

52. A method of inhibiting the binding of a neurokinin to a neurokinin 3 receptor, said method comprising contacting a compound of claim 1 with cells expressing such a receptor in the presence of a neurokinin, wherein the compound is present at a concentration sufficient to inhibit neurokinin binding to cells expressing a cloned human NK-3 receptor *in vitro*.

53. A method for altering the signal-transducing activity of neurokinin-3 receptors, said method comprising exposing cells expressing such receptors to a compound according to claim 1 at a concentration sufficient to inhibit neurokinin binding to cells expressing a cloned human neurokinin-3 receptor *in vitro*.

54. A packaged pharmaceutical composition comprising the pharmaceutical composition of Claim 23 in a container and instructions for using the composition to treat a patient suffering from a disorder responsive to neurokinin-3 receptor antagonism.

55. The packaged pharmaceutical composition of claim 35, wherein said patient is suffering from anxiety, depression, schizophrenia, obesity, chronic pulmonary obstructive disorder, or pain.

56. A compound according to claim 1 wherein in a standard assay of neurokinin-3 receptor binding the compound exhibits an IC_{50} of 1 micromolar or less.

57. A compound according to claim 1 wherein in a standard assay of neurokinin-3 receptor binding the compound exhibits an IC_{50} of 100 nanomolar or less.

58. A compound according to claim 1 wherein in a standard assay of neurokinin-3 receptor binding the compound exhibits an IC_{50} of 10 nanomolar or less.

5

INTERNATIONAL SEARCH REPORT

Inter national Application No

PCT/US 00/11187

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D215/50 C07D401/04 A61K31/4709 A61P25/00 A61P25/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, EPO-Internal, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98 52942 A (SMITHKLINE BEECHAM S. P. A.) 26 November 1998 (1998-11-26) the whole document ---	1-58
A	WO 97 21680 A (SMITHKLINE BEECHAM S. P. A.) 19 June 1997 (1997-06-19) the whole document ---	1-58
A	WO 97 19928 A (SMITHKLINE BEECHAM S. P. A.) 5 June 1997 (1997-06-05) the whole document ---	1-58
A	WO 97 19926 A (SMITHKLINE BEECHAM S. P. A.) 5 June 1997 (1997-06-05) the whole document ---	1-58
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Date of the actual completion of the international search

30 August 2000

Date of mailing of the international search report

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Kyriakakou, G

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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A	WO 96 02509 A (SMITHKLINE BEECHAM FARMACEUTICI S. P. A.) 1 February 1996 (1996-02-01) the whole document ----	1-58
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